



# Exergy Based Microturbine Analysis

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- What is Exergy?
- First Law Versus Second Law

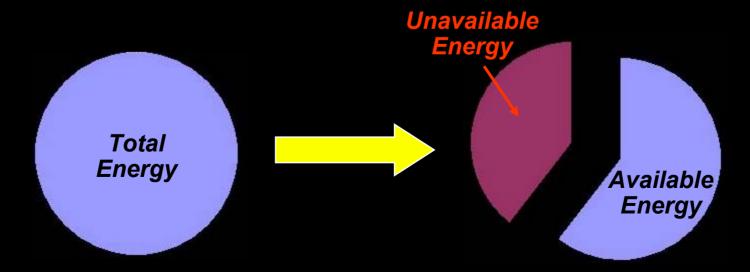


- Second Law Application Air Standard Brayton Cycle
- Chemical Availability And Combustion Losses
- Summary



#### What is Exergy?

- Exergy Represents the Maximum Work Potential A Device Can Deliver
- Term Made Popular in Europe in 1950's
- Synonymous with "Availability" (Term introduced by M.I.T. in 1940's)
- Unlike Energy, Exergy Can Be Destroyed
  - ✓ Friction
  - ✓ Uncontrolled Expansion/Compression
  - ✓ Heat Transfer
- Allows For Greater Understanding of Component and System Efficiencies





#### First Law of Thermodynamics

$$M_{air\_earth} = 1.03x10^{19} lbm$$

$$T_0 = 59 \, ^{\circ}F$$

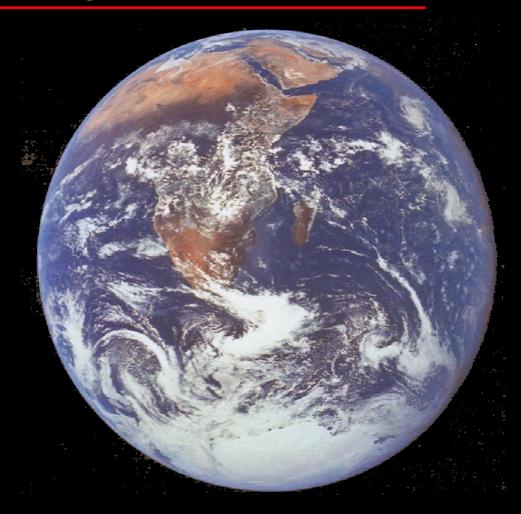
$$E_{air\ earth} = 1.278 \times 10^{21} BTU$$

$$t = 1.0 \times 10^6 \text{ years}$$

$$\eta = 0.01$$

$$P_{air\_earth} = (E_{air\_earth} \times \eta) / t$$

$$P_{air\_earth} = 427.4 \text{ MW}$$



The First Law of Thermodynamics deals with the *quantity* of energy and asserts that energy cannot be created or destroyed



## Second Law of Thermodynamics

$$M_{air earth} = 1.03x10^{19} lbm$$

$$T_0 = 59 \, ^{\circ}F$$

$$E_{air earth} = 1.278 \times 10^{21} BTU$$

$$t = 1.0x10^6 \text{ years}$$

$$\eta = 0.01$$

$$W_{air\_earth} = (1-T_{sink}/T_{source}) \times Q_{in}$$

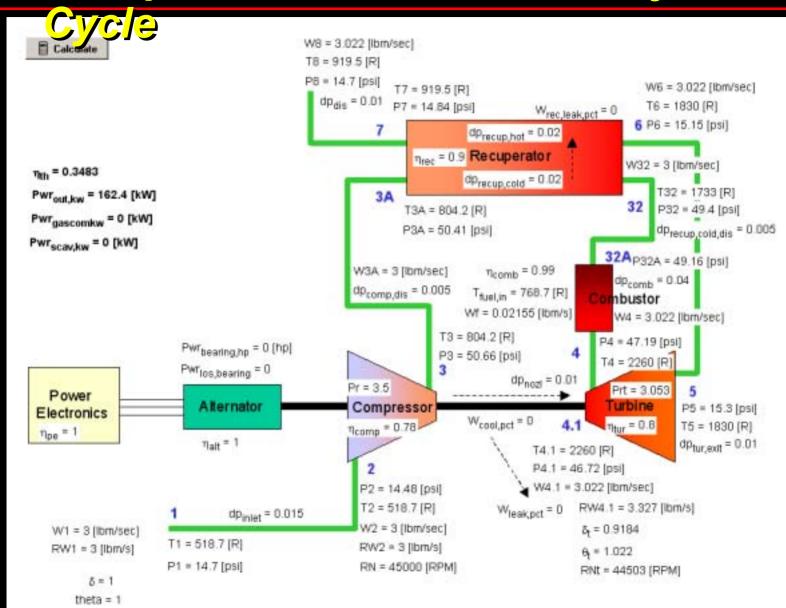
$$W_{air\_earth} = 0$$



The Second Law of Thermodynamics deals with the *quality* of energy and the degradation of energy during a process



#### Recuperated Air Standard Brayton



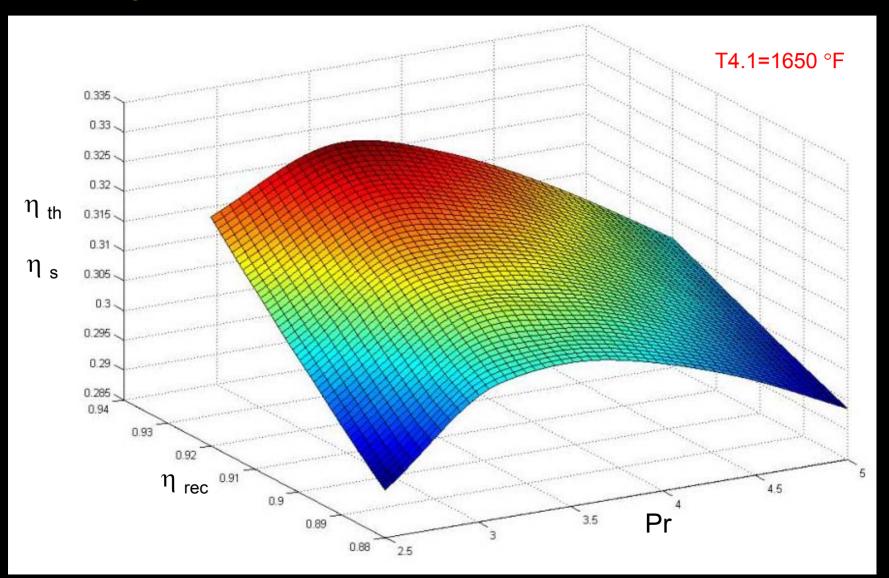


#### First And Second Law Work Together

$$q_{avail\_out} = (h_8 - h_1) - (s_8 - s_1) \cdot T_0$$
 
$$\eta_{rec} = \frac{h_{32} - h_{3a}}{h_6 - h_{3a}}$$
 
$$\eta_{th} = \frac{P_{out}}{W_{f'}LHV_{CH4}}$$
 
$$W_{6} \cdot h_6 + W_{3a} \cdot h_{3a} = W_{32} \cdot h_{32} + W_{7} \cdot h_7 + Q$$
 
$$W_{av} = Q_{in\_total} - q_{avail\_out} - \Delta S_{total} \cdot T_0$$
 
$$\eta_{rec} = 0.90$$
 
$$\eta_{s} = \frac{W_{av}}{Q_{in\_total}}$$
 
$$q_{in\_total} - q_{avail\_out} - \Delta S_{total} \cdot T_0$$
 
$$q_{in\_total} - q_{avail\_out} - \Delta S_{total} - q_{avail\_out} - \Delta S_{total} \cdot T_0$$
 
$$q_{in\_total} - q_{avail\_out} - q_{avail\_out} - \Delta S_{total} - q_{avail\_out} - q_{a$$

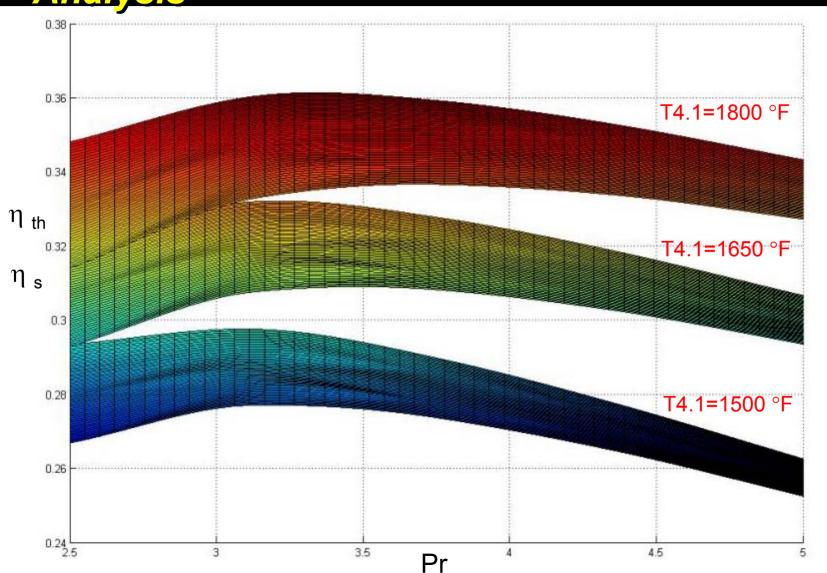


# Agreement Between First and Second Law Analysis



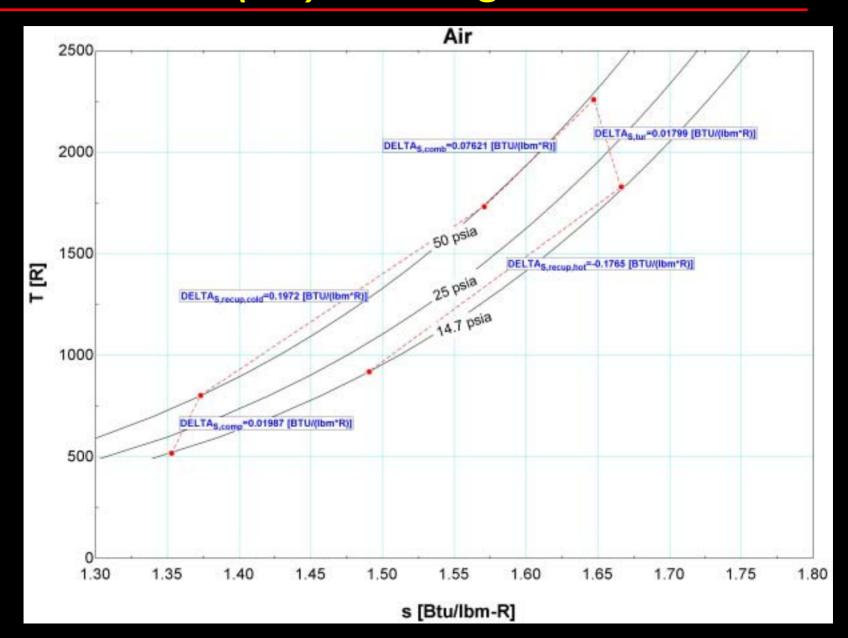


Agreement Between First and Second Law Analysis



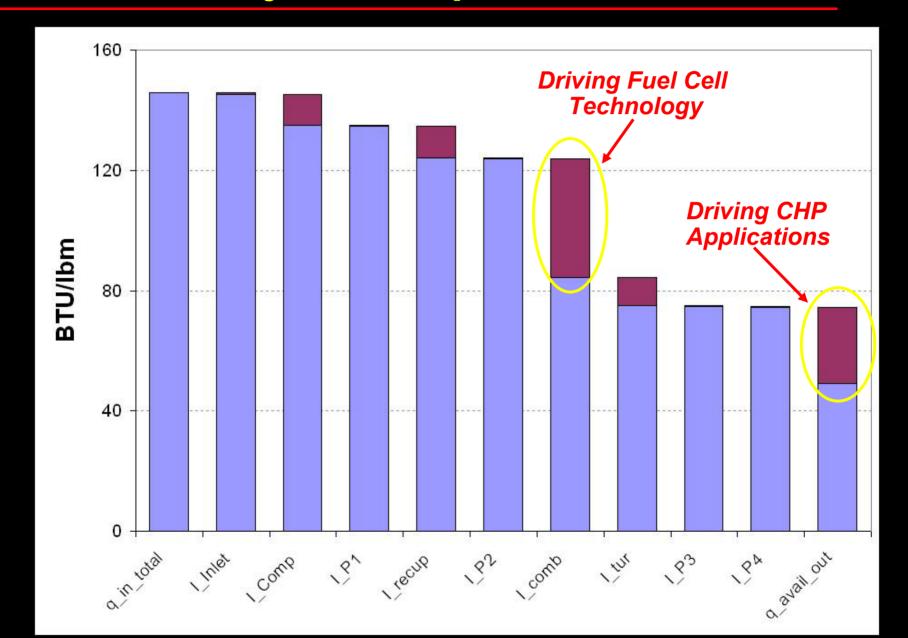


#### Ideal Gas (Air) T-s Diagram





## Availability Stack-up





#### **Calculating Combustion Losses**

CH4 + 3 (O2+3.76N2) = 0.9996 C02 + 0.00036 CO + 2 H2O + 11.26 N2 + 0.983 O2 + 0.03442 NO

$$\sum \left[ N_p \cdot \left( h_{fp}^{\ o} + h_p^{\ o} \right) \right] = \sum \left[ N_r \cdot \left( h_{fr}^{\ o} + h_r^{\ o} \right) \right] = \left( N \cdot h_f^{\ o} \right)_{CH4}$$

$$\Phi = 0.6667$$
  $T_{prod} = 1784K$ 

$$\Delta S_{\text{surr}} = 0$$

$$S_{gen} = \Delta S_{sys} = S_{prod} - S_{react} = \sum N_p \cdot S_p - \sum N_r \cdot S_r$$

$$S_X = N_X \cdot S_X(T, P_X) = N_X \cdot \left[ S_X^o(T, P_0) - R \cdot ln(y_X \cdot P_{total}) \right]$$

$$I = T_o \cdot S_{gen}$$

$$I = 288350 \frac{kJ}{\text{kmol K}}$$





# Calculating Combustion Losses

CH4 + 3 (O2+3.76N2) = 0.9996 C02 + 0.00036 CO + 2 H2O + 11.26 N2 + 0.983 O2 + 0.03442 NO

$$\frac{P_{\text{sat}}}{P_{\text{ref}}} \cdot (4.76 \text{ N}_{\text{O2}} + \text{N}_{\text{V}}) = \text{N}_{\text{V}}$$
  $N_{\text{V}} = 0.463 \text{ kmol}$   $N_{\text{I}} = 1.537 \text{ kmol}$ 

$$N_V = 0.463 \,\mathrm{kmol}$$

$$N_1 = 1.537 \cdot kmol$$

$$Q = \sum \left[ N_{p} \cdot \left( h_{fp}^{o} + h_{p}^{o} \right) \right] - \sum \left[ N_{r} \cdot \left( h_{fr}^{o} + h_{r}^{o} \right) \right] \qquad T_{prod} = T_{o}$$

$$T_{prod} = T_{o}$$

$$Q = -866686 \frac{kJ}{kmol}$$

$$\Delta S_{\text{sys}} = S_{\text{prod}} - S_{\text{react}}$$

$$\Delta S_{\text{sys}} = S_{\text{prod}} - S_{\text{react}}$$
  $S = \sum_{i} N_{x} \left[ s_{x}^{o} (T_{o}, P_{o}) - R \cdot \ln(y_{x} \cdot P_{\text{total}}) \right]$ 

$$\Delta S_{surr} = \frac{-Q_{surr}}{T_{o}}$$

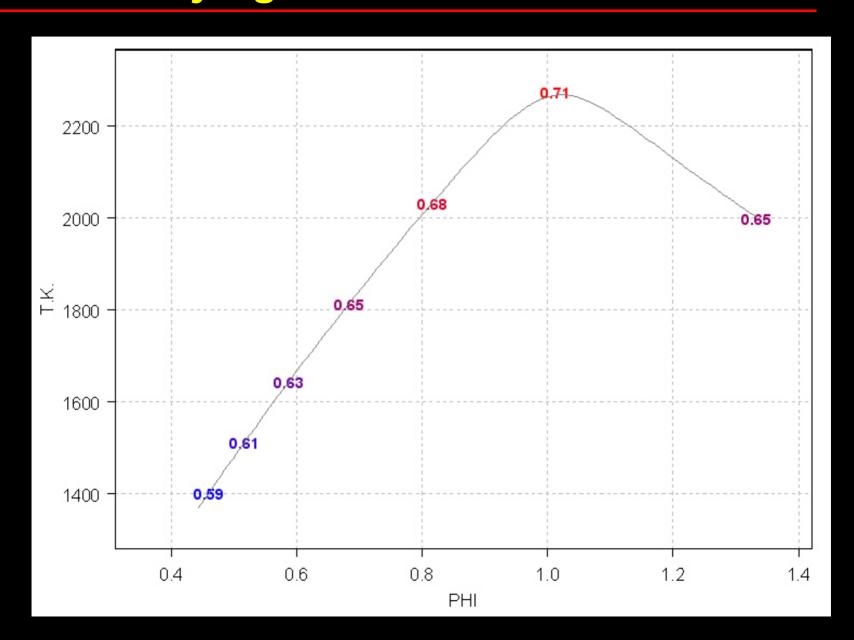
$$\Delta S_{\text{surr}} = \frac{-Q_{\text{surr}}}{T_{\text{o}}}$$
  $S_{\text{gen}} = \Delta S_{\text{sys}} + \Delta S_{\text{surr}} = 2769 \frac{\text{kJ}}{\text{kmol}_{\text{CH4}}}$ 

$$I = T_o \cdot S_{gen} = 825268 \frac{kJ}{kmol_{CH4}}$$

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  $\eta_S = \frac{I - W_{irr}}{I} = \frac{825268 - 288350}{825268} = 0.650$ 



#### **Quantifying Combustion Losses**





#### Second Law Summary

- Determines the Maximum Work Potential of Any Thermal/Fluid System
- Provides Greater Insight To System Losses Than A First Law Analysis Alone
- A Powerful Tool To Optimize Thermal Efficiency
- Is No More Difficult To Understand or Implement Than First Law Analysis

